# Your Guide to Understanding Genetic Conditions

# PHF8 gene

PHD finger protein 8

#### **Normal Function**

The *PHF8* gene provides instructions for making a protein that is found in the cell nucleus, particularly in brain cells before and just after birth. The PHF8 protein is part of a group known as zinc finger proteins, which contain one or more short regions called zinc finger domains. These regions include a specific pattern of protein building blocks (amino acids) and one or more charged atoms of zinc (zinc ions). The folded configuration of the zinc finger domain stabilizes the protein and allows it to attach (bind) to other molecules.

The PHF8 protein contains a specific zinc finger domain called a PHD domain, which binds to complexes called chromatin, the network of DNA and proteins (called histones) that packages DNA into chromosomes. Binding with the PHF8 protein is part of the process that changes the structure of chromatin (chromatin remodeling) to alter how tightly regions of DNA are packaged. Chromatin remodeling is one way gene activity (expression) is regulated; when DNA is tightly packed genes tend to be turned off, compared to when DNA is loosely packed and genes are usually turned on. While the PHF8 protein is bound to chromatin, another domain of the PHF8 protein, called Jumonji C (JmjC), removes molecules called methyl groups from histones. Removing these methyl groups (demethylation) causes the chromatin to become loosely packed and increases the expression of specific genes.

## **Health Conditions Related to Genetic Changes**

X-linked intellectual disability, Siderius type

At least four mutations in the *PHF8* gene have been found to cause X-linked intellectual disability, Siderius type. This condition is characterized by mild to moderate intellectual disability, and it occurs only in males. Affected boys often have an opening in the lip (cleft lip) with an opening in the roof of the mouth (cleft palate). Most *PHF8* gene mutations lead to an abnormally short protein that gets transported out of the cell's nucleus. Outside of the nucleus, the PHF8 protein cannot interact with chromatin to regulate gene expression. Other mutations impair the protein's ability to remove methyl groups from histones within chromatin, leading to a decrease in gene expression.

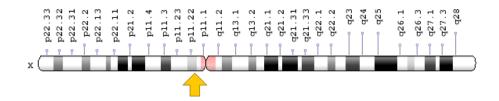
While the exact disease mechanism is unknown, it is likely that impaired protein function or a lack of PHF8 protein in the nucleus of brain cells before birth prevents chromatin remodeling, altering the normal expression of genes involved in intellectual

function and formation of structures along the midline of the skull. This altered gene expression leads to intellectual disability and cleft lip and palate found in males with X-linked intellectual disability, Siderius type.

#### **Chromosomal Location**

Cytogenetic Location: Xp11.22, which is the short (p) arm of the X chromosome at position 11.22

Molecular Location: base pairs 53,936,680 to 54,048,935 on the X chromosome (Homo sapiens Annotation Release 108, GRCh38.p7) (NCBI)



Credit: Genome Decoration Page/NCBI

#### Other Names for This Gene

- histone lysine demethylase PHF8
- JHDM1F
- jumonji C domain-containing histone demethylase 1F
- KIAA1111
- PHF8\_HUMAN
- ZNF422

#### **Additional Information & Resources**

Scientific Articles on PubMed

 PubMed https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28PHF8%5BTIAB%5D%29+OR +%28PHD+finger+protein+8%5BTIAB%5D%29%29+AND+%28%28Genes %5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND +english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+3600+days %22%5Bdp%5D

#### **OMIM**

 PHD FINGER PROTEIN 8 http://omim.org/entry/300560

#### Research Resources

- Atlas of Genetics and Cytogenetics in Oncology and Haematology http://atlasgeneticsoncology.org/Genes/GC PHF8.html
- Biochemistry (fifth edition, 2002): The Control of Gene Expression Requires Chromatin Remodeling https://www.ncbi.nlm.nih.gov/books/NBK22479/#A4460
- ClinVar https://www.ncbi.nlm.nih.gov/clinvar?term=PHF8%5Bgene%5D
- HGNC Gene Family: Lysine demethylases http://www.genenames.org/cgi-bin/genefamilies/set/485
- HGNC Gene Family: PHD finger proteins http://www.genenames.org/cgi-bin/genefamilies/set/88
- HGNC Gene Symbol Report http://www.genenames.org/cgi-bin/gene\_symbol\_report?q=data/ hgnc\_data.php&hgnc\_id=20672
- NCBI Gene https://www.ncbi.nlm.nih.gov/gene/23133
- UniProt http://www.uniprot.org/uniprot/Q9UPP1

# **Sources for This Summary**

- Fortschegger K, de Graaf P, Outchkourov NS, van Schaik FM, Timmers HT, Shiekhattar R. PHF8 targets histone methylation and RNA polymerase II to activate transcription. Mol Cell Biol. 2010 Jul; 30(13):3286-98. doi: 10.1128/MCB.01520-09. Epub 2010 Apr 26.
   Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20421419
   Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2897584/
- Kleine-Kohlbrecher D, Christensen J, Vandamme J, Abarrategui I, Bak M, Tommerup N, Shi X, Gozani O, Rappsilber J, Salcini AE, Helin K. A functional link between the histone demethylase PHF8 and the transcription factor ZNF711 in X-linked mental retardation. Mol Cell. 2010 Apr 23; 38(2):165-78. doi: 10.1016/j.molcel.2010.03.002. Epub 2010 Mar 25.
   Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20346720
   Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2989439/

• Loenarz C, Ge W, Coleman ML, Rose NR, Cooper CD, Klose RJ, Ratcliffe PJ, Schofield CJ. PHF8, a gene associated with cleft lip/palate and mental retardation, encodes for an Nepsilon-dimethyl lysine demethylase. Hum Mol Genet. 2010 Jan 15;19(2):217-22. doi: 10.1093/hmg/ddp480. Epub 2009 Oct 19.

Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/19843542
Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4673897/

- OMIM: PHD FINGER PROTEIN 8 http://omim.org/entry/300560
- Qiu J, Shi G, Jia Y, Li J, Wu M, Li J, Dong S, Wong J. The X-linked mental retardation gene PHF8 is a histone demethylase involved in neuronal differentiation. Cell Res. 2010 Aug;20(8):908-18. doi: 10.1038/cr.2010.81. Epub 2010 Jun 15.

Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20548336

• Yu L, Wang Y, Huang S, Wang J, Deng Z, Zhang Q, Wu W, Zhang X, Liu Z, Gong W, Chen Z. Structural insights into a novel histone demethylase PHF8. Cell Res. 2010 Feb;20(2):166-73. doi: 10.1038/cr.2010.8. Epub 2010 Jan 26.

Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20101266

### Reprinted from Genetics Home Reference:

https://ghr.nlm.nih.gov/gene/PHF8

Reviewed: June 2015 Published: March 21, 2017

Lister Hill National Center for Biomedical Communications U.S. National Library of Medicine National Institutes of Health Department of Health & Human Services